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 NEWS 18 SEP 11 CA/CAplus enhanced with more pre-1907 records
             JUNE 30 CURRENT WINDOWS VERSION IS V8.01b, CURRENT
 NEWS EXPRESS
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 26 JUNE 2006.
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=> FILE REG

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SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

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STRUCTURE FILE UPDATES: '14 SEP 2006 HIGHEST RN 906714-10-1 DICTIONARY FILE UPDATES: 14 SEP 2006 HIGHEST RN 906714-10-1

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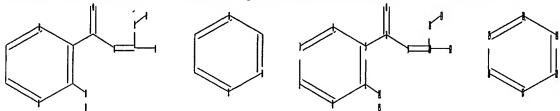
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Uploading C:\Program Files\Stnexp\Queries\SODIUM CHANNEL PYRAZINE DIV - 1.str



chain nodes :

7 8 9 10 11 12 13 14 15

ring nodes :

1 2 3 4 5 6 17 18 19 20 21 22

chain bonds :

5-7 6-14 7-8 7-9 9-10 10-11 10-12 11-13 14-15

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 17-18 17-22 18-19 19-20 20-21 21-22

exact/norm bonds :

6-14 7-8 7-9 9-10 10-11 10-12

exact bonds :

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normalized bonds :

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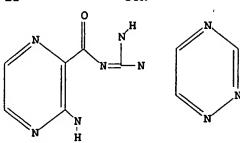
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containing 1 : 17 :

Match level :

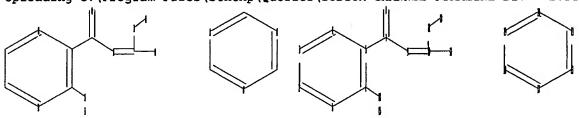
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:CLASS 9:CLASS 10:CLASS 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 17:Atom 18:Atom 19:Atom 20:Atom 21:Atom 22:Atom

=> D L1 L1 HAS NO ANSWERS L1 STR



Structure attributes must be viewed using STN Express query preparation.

=> Uploading C:\Program Files\Stnexp\Queries\SODIUM CHANNEL PYRAZINE DIV - 2.str



chain nodes :

7 8 9 10 11 12 13 14 15

ring nodes :

1 2 3 4 5 6 17 18 19 20 21 22

chain bonds :

5-7 6-14 7-8 7-9 9-10 10-11 10-12 11-13 14-15

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6 17-18 17-22 18-19 19-20 20-21 21-22

exact/norm bonds :

6-14 7-8 7-9 9-10 10-11 10-12

exact bonds :

5-7 11-13 14-15

normalized bonds :

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isolated ring systems :

containing 1 : 17 :

Match level :

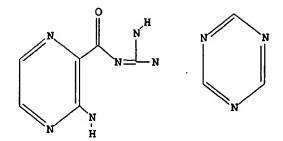
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L2 STRUCTURE UPLOADED

=> D L2

L2 HAS NO ANSWERS

L2 STR



Structure attributes must be viewed using STN Express query preparation.

=> S L1

SAMPLE SEARCH INITIATED 10:40:01 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED · 0 İTERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 0 TO 0 PROJECTED ANSWERS: 0 TO 0

L3 0 SEA SSS SAM L1

=> S L2

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SAMPLE SCREEN SEARCH COMPLETED - 2 TO ITERATE

100.0% PROCESSED 2 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE 'PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 2 TO 124
PROJECTED ANSWERS: 0 TO 0

L4 0 SEA SSS SAM L2

=> S L1 SSS FULL

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FULL SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

L5 0 SEA SSS FUL L1

=> S L2 SSS FULL

FULL SEARCH INITIATED 10:40:13 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 98 TO ITERATE

100.0% PROCESSED 98 ITERATIONS 2 ANSWERS

SEARCH TIME: 00.00.01

L6 2 SEA SSS FUL L2

=> FILE CAPLUS

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ENTRY SESSION 333.88 334.09

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=> S L6

L7 2 L6

=> D 1-2 IBIB ABS HITSTR

L7 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:346797 CAPLUS

DOCUMENT NUMBER: 142:411366

TITLE: Preparation of pyridazinylcarbonyl-substituted ureas

used for reducing risk of infection from pathogens

INVENTOR(S): Johnson, Michael R.; Hopkins, Samuel E.

PATENT ASSIGNEE(S): Parion Sciences, Inc., USA

SOURCE: PCT Int. Appl., 218 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

PA'	TENT	NO.			KIN	_	DATE						NO.		D.	ATE		
WO 2005034847												20040819						
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							DE,											
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	
		LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,	
							PL,											
							TZ,											
	RW:						MW,											
							RU,											
							GR,											
							CF,											
		SN,	TD,	TG														
US	US 2005090505				A1		2005	0428	US 2004-920626						20040818			
AU	AU 2004279329				A1		2005	0421	AU 2004-279329						20040819			
CA 2533886				AA	20050421			CA 2004-2533886						20040819				
EP 1656096				A2	20060517			EP 2004-809587						20040819				
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU.	NL.	SE,	MC,	PT,	

IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, HR 20060914 US 2005-211707 20050826 US 2006205738 A1 PRIORITY APPLN. INFO.: US 2003-496481P 20030820 US 2004-920626 20040818 20030818 US 2003-495712P P 20030818 P US 2003-495720P US 2003-495725P P 20030818 US 2004-920410 A3 20040818 WO 2004-US26963 W 20040819

OTHER SOURCE(S):

MARPAT 142:411366

GI

$$\begin{array}{c|c}
X & O & HNR^1 & R^3 \\
N = C - N & R^4 & R^4 & R^4
\end{array}$$

AB Title compds. I [X = H, halo, CF3, etc.; Y = H, OH, SH, etc.; R1 = H, alkyl; R2 = alkoxy, etc.; R3-4 = H, alkyl, OH, alkyl, Ph, etc.] are prepared For instance, II is prepared in 4 steps from [4-(4-hydroxyphenyl)butyl]carbamic acid benzyl ester (preparation given), 4-bromobutyronitrile and 1-(3,5-diamino-6-chloropyrazine-2-carbonyl)-2-methylisothiourea•HI. II has EC50 = 25 nM in a sodium channel blocker assay. I are useful for prophylactic treatment to one or more members of a population at risk of exposure to or already exposed to one or more airborne pathogens, either from natural sources or from intentional release of pathogens into the environment.

IT 847200-88-8P 847200-89-9P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of pyridazinylcarbonyl-substituted ureas used for reducing risk of infection from pathogens)

RN 847200-88-8 CAPLUS

CN Pyrazinecarboxamide, 3,5-diamino-6-chloro-N-[imino[[4-[4-[2-oxo-2-(1,3,5-triazin-2-ylamino)ethoxy]phenyl]butyl]amino]methyl]- (9CI) (CA INDEX NAME)

RN 847200-89-9 CAPLUS

CN Pyrazinecarboxamide, 3,5-diamino-6-chloro-N-[[[4-[4-[2-[(4,6-diamino-1,3,5-triazin-2-y1)amino]-2-oxoethoxy]phenyl]butyl]amino]iminomethyl]- (9CI) (CA INDEX NAME)

PAGE 1-A
NH2
O NH
C-NH-C-NH-(CH2)4

O-CH2-C-NH
N

PAGE 1-B

-NH2

L7 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:177896 CAPLUS

DOCUMENT NUMBER:

142:280225

TITLE:

Preparation of capped aminopyrazinoylguanidines as

sodium channel blockers

INVENTOR(S):

Johnson, Michael R.; Molino, Bruce F.; Zhang,

Jianzhong; Sargent, Bruce J.

PATENT ASSIGNEE(S):

Parion Sciences, Inc., USA

SOURCE:

PCT Int. Appl., 100 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE			
		D1111	APPLICATION NO.	DAIL			
WO 2005018644	A1	20050303	WO 2004-US26885	20040818			
WO 2005018644	B1	20050512					
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			DM, DZ, EC, EE, EG, ES,				
			IN, IS, JP, KE, KG, KP,				
LK, LR, L	, LT, LU	, LV, MA,	MD, MG, MK, MN, MW, MX,	MZ, NA, NI,			
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AU 2004266704	A1	20050303	AU 2004-266704	20040818			
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US	2005080091				A1	2	2005	0414	τ	JS	2004-	920	110			2004	081	18
US	7064129	9			B2	2	2006	0620										
EP	1663235				A1	20060607			1	2004 -			20040818					
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PRIORITY	APPLN.	. IN	FO.	:					τ	JS :	2003-	495	725P	1	P	2003	081	L8
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									V	NO :	2004-	US26	885	ī	N	2004	081	18

OTHER SOURCE(S):

MARPAT 142:280225

I

GI

ΔR Title compds. [I; X = H, halo, CF3, alkyl, (substituted) Ph, etc.; Y = H, OH, SH, alkoxy, alkylthio, halo, alkyl, (substituted) aryl, etc.; R1 = H, alkyl; R2 = R7, (CH2)mOR8, (CH2)mNR7R10, (CH2CH2O)mR8, etc.; m = 1-7; R3, R4 = H, alkyl, hydroxyalkyl, Ph, phenylalkyl, naphthylalkyl, pyridylalkyl, etc.; R7 = H, alkyl, (substituted) Ph, etc.; R8 = H, alkyl, 2-tetrahydropyranyl, glucuronide, etc.; R10 = H, S02Me, C0R13, C02R13, etc.; R13 = H, R7, R10, etc.; with provisos], were prepared Thus, [4-(4-hydroxyphenyl)butyl]carbamic acid benzyl ester in EtOH at 70° was treated with oxiranylmethanol over 4 h to give 4.6% [4-[4-[3-(2,3-dihydroxypropoxy)-2-hydroxypropoxy]phenyl]butyl]carbamic acid benzyl ester. This was hydrogenolyzed in EtOH over Pd/C to give 51% 3-[3-[4-(4-aminobutyl)phenoxy]-2-hydroxypropoxy]propane-1,2-diol. The latter was stirred with Et3N and 1-(3,5-diamino-6-chloropyrazine-2carbonyl)-2-methylisothiourea hydroiodide in EtOH at 65° to give 36% N-(3,5-diamino-6-chloropyrazine-2-carbonyl)-N'-[4-[4-[3-(2,3dihydroxypropoxy)-2-hydroxypropoxy]phenyl]butyl]guanidine (PSA 15143). The latter showed Na channel blocking activity with EC50 = 7 nM. TT 847200-88-8P 847200-89-9P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(claimed compound; preparation of aminopyrazinoylguanidines as sodium channel

blockers)

RN 847200-88-8 CAPLUS

CN Pyrazinecarboxamide, 3,5-diamino-6-chloro-N-[imino[[4-[4-[2-oxo-2-(1,3,5-triazin-2-ylamino)ethoxy]phenyl]butyl]amino]methyl]- (9CI) (CA INDEX NAME)

RN 847200-89-9 CAPLUS

CN Pyrazinecarboxamide, 3,5-diamino-6-chloro-N-[[[4-[4-[2-[(4,6-diamino-1,3,5-triazin-2-yl)amino]-2-oxoethoxy]phenyl]butyl]amino]iminomethyl]- (9CI) (CA INDEX NAME)

PAGE 1-B

 \sim NH₂

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> LOG HOLD COST IN U.S. DOLLARS SINCE FILE TOTAL **ENTRY** SESSION FULL ESTIMATED COST 10.68 344.77 DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE -1.50 -1.50

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